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# Pharmacokinetics and pharmacodynamics of orally administered acetylenic tricyclic *bis*(cyanoenone), a highly potent Nrf2 activator with a reversible covalent mode of action

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## ABSTRACT

The acetylenic tricyclic *bis*(cyanoenone) TBE-31 is a highly potent cysteine targeting compound with a reversible covalent mode of action; its best-characterized target being Kelch-like ECH-associated protein-1 (Keap1), the cellular sensor for oxidants and electrophiles. TBE-31 reacts with cysteines of Keap1, impairing its ability to target nuclear factor-erythroid 2 p45-related factor 2 (Nrf2) for degradation. Consequently, Nrf2 accumulates and orchestrates cytoprotective gene expression. In this study we investigated the pharmacokinetic and pharmacodynamic properties of TBE-31 in C57BL/6 mice. After a single oral dose of 10  $\mu\text{mol/kg}$  ( $\sim 200$  nmol/animal), the concentration of TBE-31 in blood exhibited two peaks, at 22.3 nM and at 15.5 nM, 40 min and 4 h after dosing, respectively, as determined by a quantitative stable isotope dilution LC-MS/MS method. The  $\text{AUC}_{0-24\text{h}}$  was 195.5 h/nmol/l, the terminal elimination half-life was 10.2 h, and the  $k_{\text{el}}$  was  $0.068\text{ h}^{-1}$ . To assess the pharmacodynamics of Nrf2 activation by TBE-31, we determined the enzyme activity of its prototypic target, NAD(P)H:quinone oxidoreductase 1 (NQO1) and found it elevated by 2.4- and 1.5-fold in liver and heart, respectively. Continuous feeding for 18 days with diet delivering the same daily doses of TBE-31 under conditions of concurrent treatment with the immunosuppressive agent azathioprine had a similar effect on Nrf2 activation without any indications of toxicity. Together with previous reports showing the cytoprotective effects of TBE-31 in animal models of carcinogenesis, our results demonstrate the high potency, efficacy and suitability for chronic administration of cysteine targeting reversible covalent drugs.

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## 1. Introduction

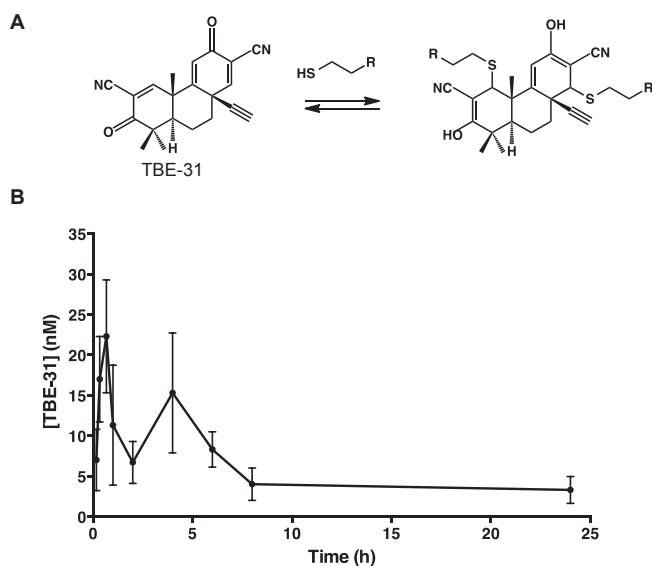
Electrophilic olefins react readily with nucleophilic cysteine sulfhydryl groups in a reversible manner. The presence of an electron withdrawing nitrile group adjacent to an olefin increases the reactivity, and furthermore, enhances the reversibility of the reaction [1]. The ability of small molecules to bind their protein targets covalently but reversibly is gaining an increasing interest in drug development as such compounds provide sustained covalent

inhibition without accumulating permanently modified (and potentially cytotoxic) proteins. The concept of designing reversible covalent inhibitors has been recently employed to obtain non-catalytic cysteine targeting drug candidates with high ligand efficiency and selectivity for the MSK/RSK-family protein kinases [2]. A reversible covalent mode of action combines the desirable features of both irreversible covalent agents (such as high potency and long-lasting activity) as well as reversible non-covalent drugs (such as lack of permanent target modification). One example of a cysteine targeting compound with a reversible covalent mode of action is the acetylenic tricyclic *bis*(cyanoenone) ( $\pm$ )-(4bS,8aR,10aS)-10a-ethynyl-4b,8,8-trimethyl-3,7-dioxo-3,4b,7,8,8a,9,10, 10a-octahydrophenanthrene-2,6-dicarbonitrile (TBE-31, Fig. 1A).

TBE-31 has two doubly activated Michael acceptors that render

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**Fig. 1.** (A) TBE-31 binds to sulfhydryl groups in a Michael addition reaction forming covalent, but reversible adducts. (B) Pharmacokinetics of TBE-31 in female C57/BL6 mice ( $n = 3$ ). TBE-31 (10  $\mu\text{mol/kg}$ ) was administered by oral gavage. Blood (10  $\mu\text{l}$ ) was drawn from the tail vein at 10, 20, 40, 60 min post-dosing, and subsequently at 2, 4, 6, 8, and 24 h. The blood levels of TBE-31 were determined at each time point using a stable isotope dilution LC-MS method on nano-flow LC-Orbitrap. Values are means  $\pm$  1 S.E.M.

this compound highly reactive with sulfhydryl groups (Fig. 1A). The best-characterized intracellular target of TBE-31 is Kelch-like ECH-associated protein-1 (Keap1), the protein sensor for oxidants and electrophiles [3]. Under homeostatic conditions, Keap1 binds transcription factor nuclear factor-erythroid 2 p45-related factor 2 (Nrf2) [4,5] and functions as a substrate adapter for the Cul3-mediated ubiquitination and proteasomal degradation of Nrf2 [6–8] by use of a highly efficient cyclic mechanism [9]. Consequently, the half-life of Nrf2 is very short ( $\sim 10$ – $20$  min) [10]. Pharmacological activators of Nrf2 react with specific cysteine residues of Keap1 [3,11–13], disrupting the cycle of Nrf2 degradation and leading to Nrf2 accumulation, nuclear translocation and enhanced target gene expression. The battery of Nrf2 transcriptional targets comprises genes encoding proteins with versatile cytoprotective functions, including antioxidant, anti-inflammatory and metabolic enzymes [14].

TBE-31 is one of the most potent Nrf2 activators known to date. In vitro, this acetylenic tricyclic bis(cyanoenone) binds reversibly to sulfhydryl groups such as those present in the reduced form of Cleland's reagent (dithiothreitol, DTT) [15,16] and to cysteine sensors of recombinant Keap1 [16,17]. In cells, at sub-to low nanomolar concentrations, TBE-31 induces Nrf2 transcriptional targets, and inhibits pro-inflammatory responses, such as the transcriptional upregulation of inducible nitric oxide synthase (iNOS) mediated by lipopolysaccharide or interferon- $\gamma$  (IFN- $\gamma$ ). In vivo, small (nmol) doses of TBE-31 robustly induce Nrf2-dependent cytoprotective responses [16,18] and protect animals against hepatocarcinogenesis induced by aflatoxin [15], as well as against cutaneous carcinogenesis mediated by solar-simulated ultraviolet radiation [19]. The aims of the present study were to: (i) define the pharmacokinetics of a single dose of orally administered TBE-31 in mice, (ii) evaluate the pharmacodynamics of Nrf2 activation by TBE-31 when administered either at a single dose or chronically, and (iii) to evaluate the safety of chronic administration of TBE-31 as exemplar of a highly potent Nrf2 activator with a reversible covalent mode of action.

## 2. Materials and methods

### 2.1. Materials

All reagents were of the highest purity available, purchased from common commercial suppliers. ( $\pm$ )-TBE-31 and a stable isotope labeled ( $\pm$ )-[ $^{13}\text{C}_2^{15}\text{N}_2$ ]-TBE-31 were synthesized as described [17,20].

### 2.2. Animals and treatments

The animal experiments were performed in accordance with the regulations described in the UK Animals (Scientific Procedures) Act 1986, and were in strict compliance with institutional guidelines. C57BL/6 mice were bred in our facility and maintained on a 12-h light/12-h dark cycle, 35% humidity with free access to water and pelleted RM1 diet (SDS Ltd., Witham, Essex, UK). All experimental animals were female, 6–12 weeks of age, uniformly distributed between the treated and the control groups.

For single dose experiments, TBE-31 [10  $\mu\text{mol/kg}$ , or  $\sim 200$  nmol/mouse ( $n = 3$ ), dissolved in 2% DMSO (v/v) in corn oil] was administered by oral gavage. Control mice received an equivalent volume of 2% DMSO (v/v) in corn oil. Blood (10  $\mu\text{l}$ ) was drawn from the tail vein at 10, 20, 40, 60 min post-dosing, and subsequently at 2, 4, 6, 8, and 24 h. The blood was collected in heparin-containing tubes and stored at  $-80^\circ\text{C}$  until analysis.

For continuous dietary administration of TBE-31, the compound was mixed with powdered RM1 diet (27.6 mg of TBE-31 per kg of food). The estimated dose of TBE-31 delivered by this diet was  $\sim 10$   $\mu\text{mol/kg}$ , or  $\sim 200$  nmol/mouse ( $n = 6$ ) per day. The TBE-31 diet was stored at  $4^\circ\text{C}$ , and the feeders were refilled daily over the course of the experiment (18 days). Control mice ( $n = 6$ ) were fed RM1 diet that was stored and delivered under identical conditions. The animals in both groups were receiving azathioprine (10 mg/kg) in the drinking water. Body weights were recorded twice a week. At termination of the experiments, the animals were euthanized by  $\text{CO}_2$  asphyxiation, and immediately exsanguinated by cardiac puncture. Blood was drawn in EDTA-containing tubes, plasma was isolated by centrifugation, and stored at  $-80^\circ\text{C}$  until analysis. Livers, hearts, skin, kidneys, and stomachs were harvested, immediately frozen in liquid  $\text{N}_2$ , and stored at  $-80^\circ\text{C}$ .

### 2.3. Determination of blood levels of TBE-31

The blood levels of TBE-31 were determined using a stable isotope dilution LC-MS method. Briefly, 10  $\mu\text{l}$  whole blood was extracted in 0.5 ml acetonitrile using 500 pmol [ $^{13}\text{C}_2^{15}\text{N}_2$ ]-TBE-31 (10  $\mu\text{l}$  of a 50  $\mu\text{M}$  solution) as an internal standard. The sample was sonicated in a water bath for 10 min, and subjected to centrifugation at  $13,000 \times g$  for 20 min at  $4^\circ\text{C}$ . The supernatant was transferred to a new tube, diluted to 10% acetonitrile (v/v), and subjected to solid phase extraction on an HLB Oasis Cartridge (30 mg, Waters, Manchester). Following a wash with 40% acetonitrile/0.1% formic acid (v/v) in water, the analytes were eluted with 70% acetonitrile (v/v) in water, dried under vacuum, and resuspended in 50  $\mu\text{l}$  of 2% acetonitrile/0.1% formic acid (v/v) in water. LC-MS/MS was performed on nano-flow LC-Orbitrap as described in detail [20]. To determine pharmacokinetic parameters, logarithmic linear regression was used to calculate the slope of the terminal disposition phase, based on which the terminal phase elimination half-life ( $t_{1/2}$ ) and the elimination constant ( $K_{el}$ ) were determined using a single compartment model with curve stripping.

### 2.4. Enzyme assays

Frozen tissue was pulverized into powder under liquid  $\text{N}_2$ .

Approximately 30 mg of powder were homogenized in an ice bath in ice-cold buffer [100 mM potassium phosphate, pH 7.4; 100 mM KCl; 0.1 mM ethylenediaminetetraacetic acid (EDTA)], and subjected to centrifugation at 4 °C (15,000 × g for 10 min). Supernatant fractions were used to determine the enzyme activities of NAD(P) H:quinone oxidoreductase 1 (NQO1) with menadione as a substrate [21], glutathione S-transferase (GST) with CDNB as a substrate [22], and the protein concentration using the bicinchoninic acid (BCA) assay (Thermo Scientific).

### 2.5. Blood tests

Blood tests were performed at the Mary Lyon Centre (MRC, Harwell, UK) on plasma samples free of hemolysis that had been isolated from mice fed continuously with TBE-31 (27.6 mg of TBE-31 per kg of food) for 18 days.

### 2.6. Statistical analysis

Values are means ± 1 S.D. or 1 S.E.M., as indicated in the figure legends. The differences between groups were determined by Students t-test using Excel (Microsoft Corp.).

## 3. Results and discussion

### 3.1. Pharmacokinetics and pharmacodynamics of a single dose of orally administered TBE-31

To determine the blood levels of TBE-31, we employed a quantitative liquid chromatography/mass spectrometry-based approach coupled with the use of a stable isotope-labeled internal standard, in which both nitrile groups of TBE-31 are labeled with  $^{13}\text{C}$  and  $^{15}\text{N}$  atoms, i.e.  $(\pm)\text{-}[\text{}^{13}\text{C}_2\text{}^{15}\text{N}_2]\text{-TBE-31}$  [20]. The high sensitivity of this method (which has an estimated on-column limit of detection of 80 fmol) enabled us to examine the pharmacokinetic profile of orally administered TBE-31 by drawing small amounts (10  $\mu\text{l}$ ) of blood from the mouse tail vein. This allowed us to determine the blood levels of the compound at 9 different time-points using three animals per group. Following a single oral dose of 10  $\mu\text{mol/kg}$ , the concentration of TBE-31 in whole blood exhibited two peaks (Fig. 1B). The first peak was at 22.3 nM 40 min after dosing. After a rapid decline, a second broader peak was observed at 15.5 nM 4 h after dosing. The reason for the presence of two peaks is unclear; however it is noteworthy that sulforaphane, another potent Nrf2 activator, which reacts reversibly with sulfhydryl groups, such as

the cysteine sensors of Keap1 [3,23–25], or reduced glutathione [26–28] shows a similar pharmacokinetic behavior in Sprague–Dawley rats [29], and could be due to accumulation of glutathione conjugates, and their subsequent deconjugation to give the parent compound.

The area under the concentration–time curve (AUC) from time zero to 24 h after dosing was 195.5 h/nmol/l, the terminal phase elimination half-life ( $t_{1/2}$ ) was 10.2 h, and the  $k_{el}$  was  $0.068\text{ h}^{-1}$ . As expected, there was no detectable TBE-31 in blood from vehicle-treated animals, confirming the absence of any endogenous sources and the specificity of the detection method. Importantly, the TBE-31 concentrations that were detected in the blood of the TBE-31-treated animals are entirely consistent with the concentrations of the compound which are typically used in cell culture experiments and have been shown to activate robustly Nrf2 and to protect against the toxicities of damaging agents, such as peroxynitrite or the combination of 6-thioguanine and ultraviolet radiation [15,16,18,30].

To assess the pharmacodynamics of Nrf2 activation by TBE-31, we determined the activity of NQO1, an enzyme encoded by a prototypic Nrf2-dependent gene, in liver and heart of the animals. In comparison with vehicle-treated animals, the specific enzyme activity of hepatic NQO1 was higher by 2.4-fold ( $n = 3$ ,  $p = 0.00003$ ) 24 h after a single orally administered dose (10  $\mu\text{mol/kg}$ , or ~200 nmol/mouse) of TBE-31 (Fig. 2A). Although to a lower degree, perhaps reflecting the much higher (by ~4-fold) levels of the enzyme in heart compared to liver, cardiac NQO1 was also induced and was 1.5-fold higher in TBE-31-treated than in vehicle-treated animals ( $n = 3$ ,  $p = 0.003$ ) (Fig. 2B).

### 3.2. Pharmacodynamics of chronically administered TBE-31

Due to the reversibility of the reaction with their protein targets, reversible covalent drugs do not permanently modify their protein targets and are suitable for chronic administration. To assess the pharmacodynamic effects of chronic TBE-31 administration, the compound was mixed with the diet and the animals were fed with this diet for 18 days, receiving continuously a daily dose of ~200 nmol TBE-31. In relation to our interest in developing pre-clinical models of cutaneous carcinogenesis under immunosuppressed conditions, during the last two weeks of the experiment, the mice were also receiving the immunosuppressive agent azathioprine (10 mg/kg) in the drinking water.

The TBE-31 treatment led to upregulation of the activity of NQO1 by 3.2-, 1.3-, 1.7-, 3.2- and 3.2-fold in liver, heart, skin, kidney,

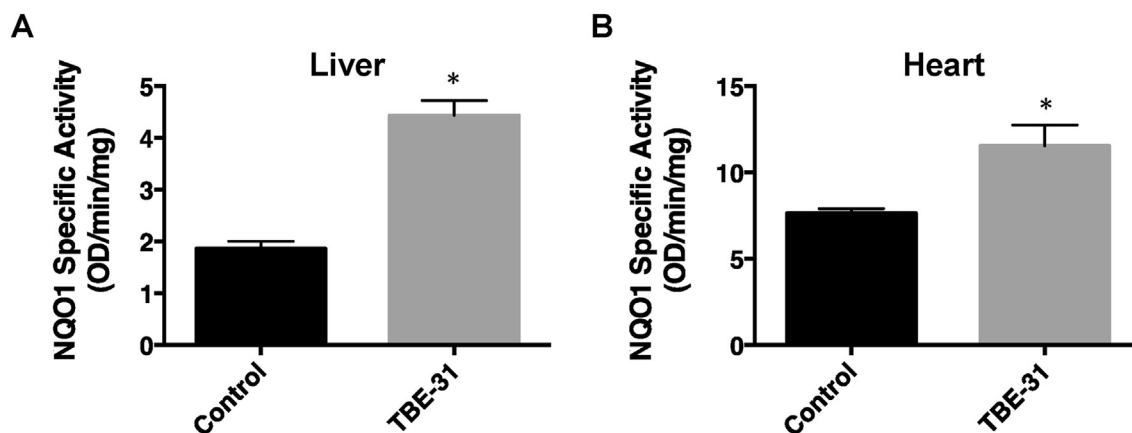
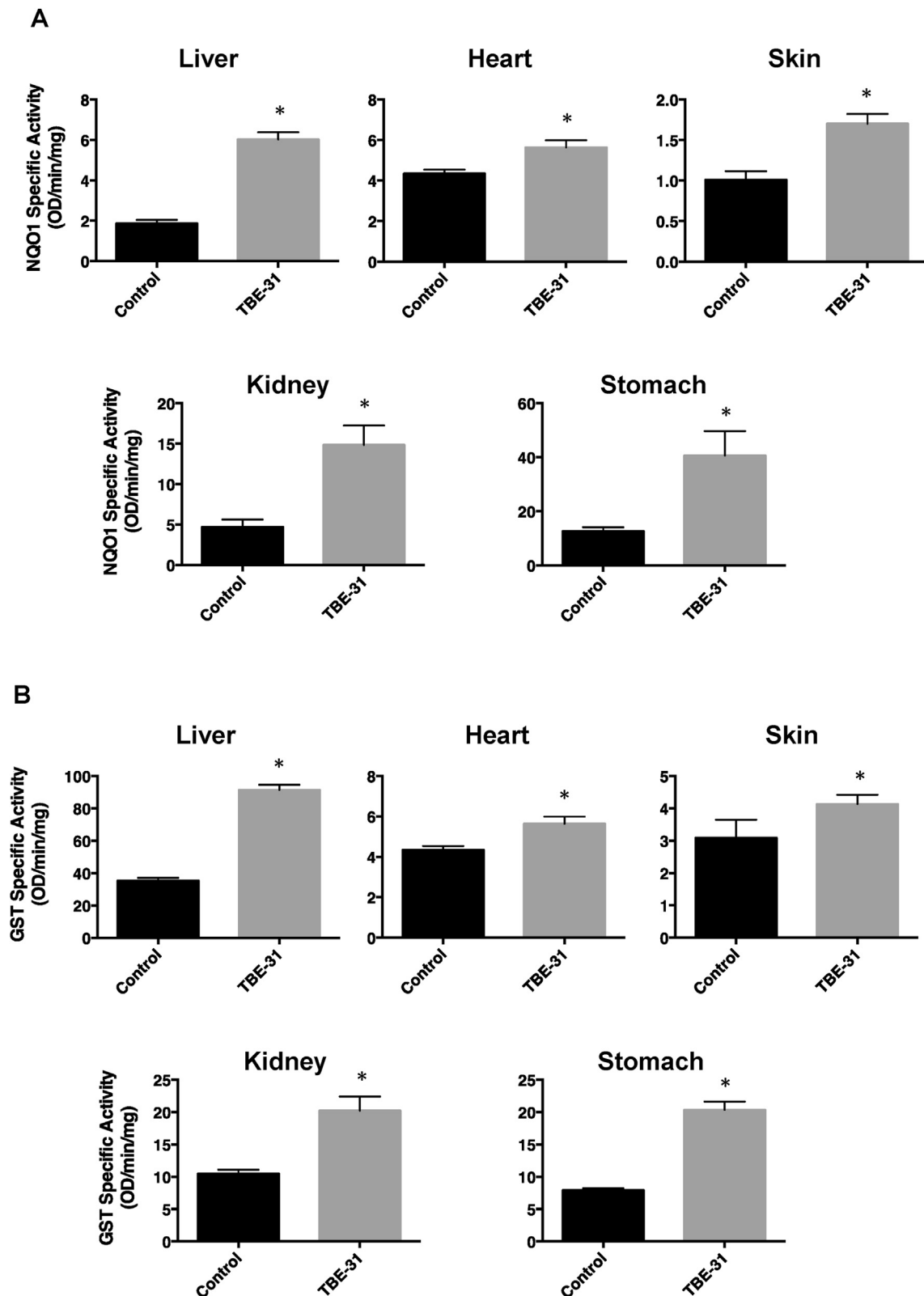


Fig. 2. Pharmacodynamics of Nrf2 activation by a single oral dose of TBE-31 in female C57/BL6 mice ( $n = 3$ ). Specific enzyme activity of NQO1 in liver (A) and heart (B) 24 h after a single oral dose (10  $\mu\text{mol/kg}$ ) of TBE-31. Values are means ± 1 S.D.

and stomach, respectively ( $n = 6$ ,  $p < 0.00001$ ) (Fig. 3A). We also measured the specific activity of glutathione *S*-transferase (GST), another Nrf2-dependent enzyme. Similar to the effect on NQO1,

chronic dietary administration of TBE-31 induced the activity of GST by 2.6-, 1.4-, 1.4-, 1.9-, and 2.6-fold in liver, heart, skin, kidney, and stomach, respectively ( $n = 6$ ,  $p < 0.00001$ ) (Fig. 3B). The highly



**Fig. 3.** Continuous feeding with TBE-31 (27.6 mg of TBE-31 per kg of food) for 18 days increases the specific enzyme activity of NQO1 (menadione as a substrate) (A) and GST (CDNB as a substrate) (B) in female C57/BL6 mice ( $n = 6$ ). Values are means  $\pm$  1 S.D.



statistically significant induction of both enzymes in all organs examined demonstrated the pharmacodynamic effects of chronic TBE-31 administration and further confirmed that TBE-31 is a potent Nrf2 activator *in vivo*.

### 3.3. Toxicity assessment of chronically administered TBE-31

There were no obvious signs of toxicity, as judged by the similarity in body weight (Fig. 4) and behavior between the animals on the control diet and those receiving continuously a daily dose of ~200 nmol TBE-31 in the diet for 18 days. To further strengthen this conclusion, we performed a detailed analysis of the plasma levels of creatinine, alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate aminotransferase (AST), total cholesterol, low-density lipoproteins (LDL), high-density lipoproteins (HDL), triglycerides, free fatty acids, glucose, bilirubin, and ketone bodies (Table 1). The activities of ALP, ALT and AST were not significantly different between the two groups, indicating unaltered liver function and confirming the absence of toxicity by the continuous TBE-31 treatment. Most of the other evaluated parameters were also very similar between the groups. There were two exceptions: the levels of LDL were higher (by 1.3-fold,  $p < 0.05$ ) in the TBE-31-treated group compared to the control group, whereas the levels of bilirubin were lower (by 35%,  $p < 0.05$ ).

The increase in LDL by the TBE-31 treatment is consistent with the pro-atherogenic effects of Nrf2 that have been previously described in apolipoprotein E (ApoE)-deficient mice [31–33]. However, it is not clear at present whether Nrf2 is pro-atherogenic in man, and perhaps more importantly, whether the pro-atherogenic role of Nrf2 requires its persistent (rather than transient) activation. Nonetheless, this result warrants caution when designing dosing regimens with Nrf2 activators in humans.

The decrease in the levels of bilirubin in the plasma of the TBE-31-treated animals is in agreement with the role of Nrf2 in regulating the gene expression of microsomal UDP-glucuronosyltransferases, such as UGT1A1 [34,35], the enzymes that are principally responsible for the glucuronidation and elimination of bilirubin [36]. This result suggests that Nrf2 activation by agents such as TBE-31 could be potentially developed as a therapeutic approach for conditions of impaired bilirubin glucuronidation, such as Crigler–Najjar and Gilbert's syndromes.

Notably, although TBE-31 is largely eliminated by 24 h after dosing, its cytoprotective effects are evident for much longer periods of time. This is because the ultimate cytoprotective agents are not TBE-31, or even Nrf2, but the Nrf2-dependent transcriptional targets, which are proteins with long half-lives. We have recently demonstrated that NQO1 is induced to essentially the same extent at 24- or 72-h after dosing with TBE-31 in the murine skin, where

**Table 1**

Blood plasma tests of samples from female C57BL/6 mice ( $n = 4–6$ ) administered continuously TBE-31 in the diet (27.6 mg of TBE-31 per kg of food) for 18 days.

Blood test	Control diet	TBE-31 diet	<i>p</i> value
Creatinine ( $\mu\text{mol/L}$ )	$8.58 \pm 0.995$	$8.75 \pm 0.704$	0.7445
Alkaline phosphatase (IU/L)	$44.0 \pm 32.5$	$16.7 \pm 9.27$	0.0810
Alanine aminotransferase (IU/L)	$49.4 \pm 26.7$	$38.3 \pm 22.5$	0.4739
Aspartate aminotransferase (IU/L)	$150.8 \pm 17.1$	$93.8 \pm 60.8$	0.2968
Total cholesterol (mmol/L)	$1.82 \pm 0.457$	$2.19 \pm 0.178$	0.2994
HDL (mmol/L)	$1.23 \pm 0.360$	$1.42 \pm 0.156$	0.6572
LDL (mmol/L)	<b><math>0.480 \pm 0.092</math></b>	<b><math>0.643 \pm 0.052</math></b>	<b>0.0150</b>
Glucose (mmol/L)	$11.6 \pm 3.05$	$15.3 \pm 1.67$	0.2211
Triglycerides (mmol/L)	$1.05 \pm 0.317$	$1.18 \pm 0.271$	0.3568
Free fatty acids (mmol/L)	$0.395 \pm 0.142$	$0.472 \pm 0.294$	0.4519
Total bilirubin ( $\mu\text{mol/L}$ )	<b><math>3.38 \pm 0.577</math></b>	<b><math>2.18 \pm 0.519</math></b>	<b>0.0478</b>
Ketone bodies (mmol/L)	$0.073 \pm 0.015$	$0.123 \pm 0.078$	0.2899

The numbers shown in bold are those that are statistically significantly different between the groups.

the half-life of the compound is ~10 h [19]. This long-lasting pharmacodynamic effect makes the maintenance of steady-state plasma levels of the inducer unnecessary, allowing for chronic dosing at a low frequency. In addition, because reversible covalent binding does not lead to formation of stable adducts with the target protein, repeated dosing regimens are possible without immunological consequences. Indeed, chronic (~30 weeks) topical application of small quantities (40 nmol per animal) of TBE-31 twice a week resulted in dramatically reduced tumor multiplicity and burden in a model of ultraviolet radiation-mediated skin carcinogenesis in SKH-1 hairless mice receiving azathioprine treatment [19]. In a rat model of aflatoxin-mediated hepatocarcinogenesis, oral administration of TBE-31, three times per week for three weeks, essentially abolished the formation of pre-neoplastic foci in liver [15]. Taken together, these findings illustrate the advantages of reversible covalent drugs and especially those whose protein targets subsequently affect the activity of transcription factors that, in turn orchestrate the expression of networks of genes, and encourage future drug development of compounds of this class.

### Conflict of interest

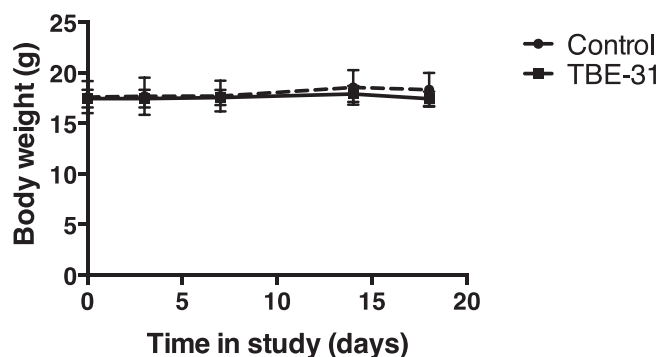
The authors declare no conflict of interest.

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**Fig. 4.** Continuous feeding with TBE-31 (27.6 mg of TBE-31 per kg of food) for 18 days does not alter body weight in female C57BL/6 mice ( $n = 6$ ). Values are means  $\pm$  1 S.D.

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